

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11) EP 1 036 553 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:

20.09.2000 Bulletin 2000/38

(51) Int. Cl.⁷: **A61K 7/02**

(21) Application number: 00105666.2

(22) Date of filing: 17.03.2000

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE

Designated Extension States: AL LT LV MK RO SI

(30) Priority: 18.03.1999 JP 7443199

(71) Applicant:

SHISEIDO COMPANY LIMITED Chuo-ku, Tokyo 104-8010 (JP) (72) Inventors:

Daisuke, Aso,
 c/o Shiseido Research Center
 Yokohama-shi, Kanagawa 223-8553 (JP)

Akihito, Yokotsuka,
 c/o Shiseido Research Center
 Yokohama-shi, Kanagawa 223-8553 (JP)

(74) Representative:

Henkel, Feiler, Hänzel Möhlstrasse 37 81675 München (DE)

(54) Solid water-in-oil type emulsion cosmetic composition

(57) A solid water-in-oil type emulsion cosmetic composition having a fresh, superior skin feeling, and effectively correcting roughness of the skin while exhibiting long-lasting power, which contains a hydrophobically treated powder, spherical silica, an oil, and water, wherein a ratio by weight of the oil to the water (i.e. oil/water) is 0.5 to 10.

Description

BACKGROUND OF THE INVENTION

Field of the Invention

[0001] The present invention relates to a solid water-in-oil type emulsion cosmetic composition. More specifically it relates to a solid water-in-oil type emulsion cosmetic composition having a superior skin feeling having a freshness, when applied to, for example, the face as a cosmetic, having a superior long-lasting power against sweat and skin oil, and capable of hiding roughness of the skin surface (pores etc.), particularly it relates to a solid water-in-oil type emulsion cosmetic composition suitable as a make-up cosmetic composition.

2. Description of the Related Art

[0002] In conventional cosmetic compositions, spherical powder is often formulated therein for the purpose of eliminating or hiding shine after application or roughness of the skin surface (specifically, pores etc.). However, in the case of emulsion cosmetic compositions of a water-in-oil type, there is the problem that a large amount of spherical powder for imparting the above effect while trying to maintain the "fresh" skin feel leads to an increase in the total amount of powder, and therefore, a higher viscosity at high temperatures, decline in fluidity, and, therefore, difficulty in filling into containers. Further, when increasing the amount of the spherical powder blended without changing the total amount of powder, the covering power or a feeling that fit on skin tends to be reduced.

SUMMARY OF THE INVENTION

[0003] The present invention was made in consideration of the above-mentioned conventional situation in the art and the object thereof is to provide a solid water-in-oil type emulsion cosmetic composition having a sufficiently satisfactory skin feel, in particular freshness, and simultaneously superior in correction of skin roughness, particularly a solid water-in-oil type emulsion cosmetic composition for a make-up cosmetic composition.

[0004] The present inventors engaged in intensive studies to solve the above-mentioned problems and, as a result, found that, by formulating a hydrophobically treated powder, spherical silica, an oil, and water in the specific ratio, it is possible to obtain a solid water-in-oil type emulsion cosmetic composition superior in freshness and other skin feeling and superior in the effect of improvement of the skin, whereby the present invention has been completed.

[0005] In accordance with the present invention, there is provided a solid water-in-oil emulsion cosmetic composition comprising: a hydrophobically treated powder, spherical silica, an oil, and water, in which the ratio by weight of the oil to the water (i.e., oil/water) is 0.5 to 10.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0006] The present invention will be explained in detail below.

[0007] The "hydrophobically treated powder" used in the present invention means those composed of a powder, which can generally be formulated into a cosmetic composition, covered with various organic compounds and may be obtained by the physical adsorption, chemical bonding, etc. to the powder. Further, the hydrophobically treated powder in the present invention means those which, when 0.5 g of the treated powder is placed in 100 ml of deionized water, and is allowed to stand at 50°C for 1 hour, the majority of the powder is found to be floating on the water surface by a visual evaluation.

[0008] The hydrophobic treatment agent for such a hydrophobically treated powder includes, a hydrophobic compound such as for example, a metal soap, oil and fat, wax, silicone compound, fluorine compound, hydrocarbon compound, fatty acid ester, lyophilic surfactant may be mentioned. In the present invention, these compounds may be used alone or in any combinations thereof. Further, as the powder to be hydrophobically treated, those explained later are suitably selected for use.

[0009] The hydrophobic treatment agents particularly preferre the present invention are silicone compounds, metal soaps, and fluorine compounds. More preferred are silicone inpounds. Especially, the use of the treated powder having a high or increased coverage density of a silicone compound with an Si-H group (preferably alkylated) on the surface thereof are preferable.

[0010] The formulating amount of the hydrophobically treated powder is 5.0 to 60.0 % by weight, preferably 10.0 to 50.0 % by weight, more preferably 15.0 to 45.0 % by weight. If the amount of the hydrophobically treated powder is less than 5.0 % by weight, the covering power, fit, etc. tend to decrease, while if more than 60.0 % by weight, the smooth spreadability at the time of application and freshness tend to be decreased and the skin feel becomes worse.

[0011] The spherical silica usable in the present invention includes any silica having a so-called spherical shape preferably having a diameter of $0.1 - 30 \mu m$, more preferably $1 - 10 \mu m$. The surface of the spherical silica may be porous or nonporous, although the use of porous spherical silica is preferable, because the nonporous silica takes an aqueous component in such a manner that the aqueous component forms a film around the spherical powder. Contrary to this, the porous spherical silica powder adsorbs the aqueous component. The formulating amount of the spherical silica is $2.0 \text{ to } 15.0 \text{ % by weight, preferably } 4.0 \text{ to } 15.0 \text{ % by weight, more preferably } 5.0 \text{ to } 10.0 \text{ % by weight, in view of, for example, the feeling of use at the time of application. If the amount of the spherical silica is less than <math>2.0 \text{ % by weight, the fillability in containers becomes worse and the smoothness after application also becomes poor. Further, if more than <math>15.0 \text{ % by weight, the decrease in a feeling that fit on skin at the time of application, a slipperiness or sliminess are observed - which are not desirable in views of skin feel.$

[0012] Examples of spherical powders, other than spherical silica, capable of formulating in the present invention are polyalkyl methacrylate powder, nylon powder, polystyrene powder, polyurethane powder, cellulose powder, cross-linkable silicone powder. Among these powders, cross-linkable silicone powder and polyalkyl methacrylate powder are particularly preferable.

[0013] The proportion of the spherical silica formulated in the total spherical powder is 30 to 100 % by weight, particularly 50 to 100 % by weight. If the proportion is less than 30 % by weight, the problem during the manufacture occurs since the filling the composition into a container becomes difficult due to the increase in the viscosity at an elevated temperature and also the problem from the functional viewpoint occurs since the correction of skin roughness with the cosmetic composition is not sufficient. In addition, when the spherical silica is formulated into the cosmetic composition, the spherical silica is preferably formulated after adsorbing an aqueous component (e.g., water, hemectant etc.) in view of the applicability and the stability. Such an adsorption can be effected simply by contacting the spherical silica with the aqueous component.

[0014] The oil formulated into the solid water-in-oil type emulsion cosmetic composition of the present invention includes any oils usually used in cosmetic compositions. For example, as liquid oils, avocado oil, tsubaki oil, macademia nut oil, mink oil, olive oil, rapeseed oil, jojoba oil, triglycerin, glycerin trioctanate, etc.; as solid oils and fats, coconut oil, hydrogenated coconut oil, palm oil, beef tallow, sheep fat, Japan wax, hydrogenated castor oil, etc.; as waxes, beeswax, candelilla wax, carnauba wax, insect wax, spermaceti, lanolin, reduced lanolin, etc.; as hydrocarbons, liquid paraffin, squalane, paraffin, ceresine, vaseline, squalene, microcrystalline wax, etc.; as higher fatty acids, lauric acid, myristic acid, palmitic acid, stearic acid, behenic acid, oleic acid, 12-hydroxystearic acid, isostearic acid, linolic acid, linoleic acid, etc.; as higher alcohols, lauryl alcohol, cetyl alcohol, stearyl alcohol, behenyl alcohol, oleyl alcohol, monostearyl glycerol ether, monopalmityl glycerol ether, cholesterol, phytosterol, isostearyl alcohol, etc.; as ester oils, isopropyl myristate, cetyl octanate, octyldodecyl myristate, butyl stearate, decyl oleate, ethyleneglycol dioctanate, diisostearyl malate, trimethylolpropane trioctanate, trimethylolpropane triisostearate, pentanerythritol tetraoctanate, glycerin trioctanate, glycerin triisostearate, ethyl acetate, butyl acetate, amyl acetate, etc.; as silicones, dimethyl polysiloxane, methylphenyl polysiloxane, methylhydrogen polysiloxane, decamethyl cyclopentasiloxane, octamethyl cyclotetrasiloxane, silicone resins forming three dimensional network structures, silicone rubber, etc. may be mentioned, but the present invention is not limited to the above-mentioned oils. Further, these oils may be used alone or in any combination thereof in the solid water-in-oil type emulsion cosmetic composition of the present invention.

[0015] The ratio of the oil to the water formulated in the solid water-in-oil type emulsion cosmetic composition is a ratio of the oil to water (i.e., oil/water) of 0.5 to 10 (ratio by weight), preferably 1 to 5. If this ratio is less than 0.5, the viscosity becomes higher at a high temperature, the fluidity is decreased, and filling in a container tends to become more difficult. If the ratio is more than 10, the skin feeling, that is, the freshness at the time of application, becomes inferior.

[0016] Further, the sum of the proportions of the oil and water formulated in the solid water-in-oil type emulsion cosmetic composition is preferably 40.0 to 90.0 % by weight. If this amount of sum is less than 40.0 % by weight, the viscosity becomes higher at a high temperature, the fluidity is decreased, and filling in a container tends to become more difficult. If the amount is more than 90.0 % by weight, it is not possible to sufficiently impart the effect of smoothness after application and correcting roughness of the skin surface.

[0017] The solid water-in-oil type emulsion cosmetic composition of the present invention may have suitably formulated it, in addition to the above components, optionally, a powder other than the hydrophobically treated powder, surfactant, lower alcohol, polyhydric alcohol, humectant, preservative, polymer other than a coating agent, antioxidant, UV blocker, fragrance, various medicines, etc. in a range of quality and quantity, which do not impair the above effects of the present invention.

[0018] As the powder capable of formulating into the cosmetic compositions of the present invention, powders usually used in cosmetic compositions may be mentioned. Examples of such powder are inorganic powders such as talc, kaolin, mica, sericite, dolomite, biotite, phlogopite, synthetic mica, vermiculite, magnesium carbonate, calcium carbonate, aluminum silicate, barium silicate, calcium silicate, magnesium silicate, metal salts of tungstic acid, magnesium, silica, zeolite, barium sulfate, sintered calcium sulfate, sintered gypsum, calcium phosphate, fluorapatite, hydroxyapatite,

ceramic powder, and metal soaps (e.g., zinc myristate, calcium palmitate, and aluminum stearate); organic powders such as polyamide resin powder, polyethylene powder, methyl polymethacrylate powder, polystyrene powder, styrene and acrylic acid copolymer resin powder, benzoguanamin resin powder, polytetrafluoroethylene powder, cellulose powder; inorganic white pigments such as titanium dioxide and zinc oxide; inorganic red pigments such as iron oxide (e.g. bengara) and iron titanate; inorganic brown pigments such as γ-iron oxide; inorganic yellow pigments such as yellow iron oxide and yellow earth; inorganic black pigments such as black iron oxide, carbon black, and lower titanium oxide; inorganic violet pigments such as mango violet and cobalt violet; inorganic green pigments such as chromium oxide, chromium hydroxide, and cobalt titanate; inorganic blue pigments such as prussian blue and ultramarine; pearl pigments such as titanium oxide coated mica, titanium oxide coated bismuth oxichloride, titanium oxide coated talc, colored titanium oxide coated mica, bismuth oxichloride, and fish scales; metal powder pigments such as aluminum powder and copper powder; lakes such as Lithol rubine BCA (Red No. 202), Lithol red (Red No. 205), Deep maroon (Red No. 220), Permatone Red (Red No. 228), Permanent red F5R (Red No. 405), Permanent orange (Orange No. 203), Benzidine Orange (Orange No. 204), Benzidine yellow G (Yellow No. 205), Hanza Yellow (Yellow No. 401), and Yellow AB (Blue No. 404); zirconium, barium, or aluminum lakes and other lakes such as Erythrosine (Red No. 3), Phloxine B (Red No. 104), Fast acid magenta (Red No. 227), Violamine R (Red No. 401), Orange II (Orange No. 205), Tartrazine (Yellow No. 4), (Yellow No. 202), Fast green FCF (Green No. 3), and Brilliant blue FCF (Blue No. 1); and natural colors such as chlorophyl and β -carotin may be mentioned.

[0019] In the solid water-in-oil type emulsion cosmetic composition of the present invention, it is possible to use a surfactant usually capable of formulating in a cosmetic compositions, without regard as to its ionic character. Specifically, as anionic surfactants, for example, soap raw materials, fatty acid soaps such as sodium laurate, higher alkyl sulfate ester salts such as sodium laurosulfate, alkyl ether sulfate ester salts such as polyoxyethylene (i.e., "POE") laurosulfate triethanol amine, N-acylsarcosine acids such as sodium lauroyl sarcosinate, higher fatty acid amide sulfonates such as sodium N-cocoyl-N-methyl taurate, phosphate ester salts such as POE stearyl ether phosphate, sulfosuccinates such as sodium di-2-ethylhexylsulfosuccinate, alkylbenzensulfonates dodecylbenzensulfonate, N-acyl glutamates such as disodium N-stearoyl glutamate, higher fatty acid ester sulfate ester salts such as sodium hydrogenated glyceryl cocoate sulfate, sulfated oils such as Turkey red oil, POE alkyl ether carboxylic acid, POE alkylaryl ether carboxylate, α-olefinsulfates, higher fatty acid ester sulfonates, secondary alcohol sulfate ester salts, higher fatty acid alkylolamide sulfate ester salts, sodium lauroyl monoethanolamide succinate, Npalmitoyl asparaginate ditriethanol amine, sodium caseine, etc. may be mentioned.

[0020] As cationic surfactants, for example, alkyl trimethyl ammonium salts such as stearyl trimethyl ammonium chloride, dialkyldimethyl ammonium salts such as distearyldimethyl ammonium chloride, alkyl pyridinium salts such as cetylpyridinium chloride, alkyl quaternary ammonium salts, alkyl dimethylbenzyl ammonium salts, alkyl isoquinolinium salts, dialkyl morphonium salts, POE alkyl amines, alkyl amine salts, polyamine fatty acid derivatives, amyl alcohol fatty acid derivatives, benzalkonium chloride, benzethonium chloride, etc. may be mentioned.

[0021] As bipolar surfactants, for example, imidazoline base bipolar surfactants such as sodium 2-undecyl-N,N,N-(hydroxyethylcarboxymethyl)-2-imidazoline, betaine base surfactants such as lauryldimethyl-aminoacetate betaine, etc. may be mentioned.

[0022] As lypophilic nonionic surfactants, for example, sorbitan fatty acid esters such as sorbitan monoisostearate and sorbitan sesquioleate, glyceryl polyglyceryl fatty acids such as glyceryl monostearate, propylene glycol fatty acid esters such as propylene glycol monostearate, hydrogenated castor oil derivatives, glyceryl alkyl ethers, dimeticone copolyol, alkyl dimeticone copolyol etc. may be mentioned

[0023] As hydrophilic nonionic surfactants, for example, POE sorbitan fatty acid esters such as POE sorbitan monostearate, POE sorbite fatty acid esters such as POE-sorbite monooleate, POE glyceryl fatty acid esters such as POE glyceryl monoisostearate, POE alkyl ethers such as POE stearyl ethers and POE cholestanol ethers, POE alkyl phenyl ethers such as POE nonyl phenyl ethers, pluaronics such as Pluronic, POE • POP alkyl ethers such as POE • polyoxy-propylene (i.e., "POP") cetyl ethers, tetra POE • tetra POP ethylene diamine condensation products such as Tetronic, POE castor oil and hydrogenated castor oil derivatives such as POE castor oil and POE hydrogenated castor oil, POE beeswax • lanolin derivatives, alkanolamides, POE propylene glycol fatty acid esters, POE alkylamines, POE fatty acid amides, sucrose fatty acid esters, POE nonylphenyl formaldehyde condensation products, alkylethoxydimethylamine-oxide, trioleylphosphoric acid etc. may be mentioned, but the invention is not limited to the above surfactants.

[0024] Further, in the solid water-in-oil emulsion cosmetic of the present invention, these surfactants may be used alone or in any combinations thereof.

[0025] The use of dimeticone copolyol, alkyl dimeticone copolyol, sorbitan fatty acid esters, POE sorbitan fatty acid esters, POE hydrogenated castor oil, glyceryl fatty acid esters, POE glyceryl fatty acid esters, etc, is especially preferable as the surfactant due to the easy filling ability to a container (i.e., preferable elevated temperature characteristics), the good feeling during the application (i.e., no stickiness), the good emulsion stability, etc.

[0026] The form which the solid water-in-oil type emulsion cosmetic composition of the present invention takes is not particularly limited. The cosmetic composition may take the form of, for example, a cosmetic base, foundation, face

powder, blusher, lipstick, mascara, eyeshadow, or eyeliner.

EXAMPLES

⁵ [0027] Preferable Examples of the present invention will be explained below, but the present invention is not limited thereto. Further, the amounts blended are shown by % by weight, unless otherwise indicated.

Examples 1 to 2 and Comparative Examples 1 to 4

[0028] Solid water-in-oil emulsion type foundations were prepared by the method explained below according to the formulations shown in the following Table 1 and Table 2 and were evaluated as to various performance as cosmetic compositions. The method of evaluation was as explained below. The results are shown in Table 3.

1. Method of Preparation

[0029] The components (1) to (6) shown in Table 1 and Table 2 were heated and mixed homogeneously at 90°C. The mixed and pulverized components (7) to (12) were then added thereto to obtain homogeneously dispersed mixture. The homogeneously melted and dispersed components (13) to (25) were then added to obtain a homogeneous emulsion. Next, the emulsion was filled in a pan, followed by cooling.

2. Method of Evaluation

[0030] The solid water-in-oil type emulsion foundation prepared was subjectively evaluated by an expert panel of 10 members based on the following standard for evaluation as to whether it had a light spreadability on the skin, freshness at time of application, smoothness after application, effect in correcting skin roughness, long-lasting power after three hours after application, and ease of fillability in a container.

(1) Standard for Evaluation of Light Spreadability on the Skin, Freshness at Time of Application, Smoothness After Application, Effect in Correcting Skin Roughness, and Long-Lasting Power after 3 Hours from Application

[0031]

15

20

30

35

- ++ (Excellent):At least nine members of panel felt effects.
- + (Good): Seven to eight members of panel felt effects.
- ± (Fair): Four to six members of panel felt effects.
- (Poor): Three or less members of panel felt effects.

(2) Standard for Evaluation of Fillability in Container

40 [0032]

- ++ (Excellent): No problem in fluidity when pouring and smooth surface after filling.
- + (Good): Shaking or other procedure required when filling to obtain smooth surface.
- \pm (Fair): Cannot fill smoothly even with shaking or other procedure.
- (Poor): No fluidity and pan cannot be filled.

50

45

Table 1

Example	1	2	3
(1) Decamethylcyclopentasiloxane		28.9	28.9
(2) Dimethyl polysiloxane	5.0		5.0
(3) Liquid paraffin	3.0		3.0
(4) 2-Ethylhexyl p-methoxycinnamate		1.5	1.5
(5) Dimeticone copolyol ¹	2.0	2.0	2.0
(6) Sorbitan sesquiisostearate	1.0		1.0
(7) Isostearic acid	0.5	0.5	0.5
(8) Solid paraffin	3.0	3.0	3.0
(9) Carnauba wax	1.0	1.0	1.0
(10) Fragrance	0.1	0.1	0.1
(11) Silicone-treated titanium oxide	12.0	12.0	12.0
(12) Titanium oxide	i –	-	1 - 1
(13) Silicone-treated iron oxide	3.0	3.0	3.0
(14) Iron oxide	-	-	- 1
(15) Iron oxide-coated titanated mica	0.5		0.5
(16) Silicone-treated sericite	3.0	3.0	3.0
(17) Sericite	-	-	-
(18) Silicone-treated talc	2.5	2.5	2.5
(19) Talc	-	-	-
(20) Porous spherical silica (particle	10.0	4.0	-
size: $4 - 6 \mu m$)		l .	1
(21) Nonporous spherical silica	-	-	4.0
(particle size: 4 - 6 μm)	}		
(22) Spherical polymethyl methacrylate	-	6.0	6.0
(23) Purified water	30.0		17.6
(24) 1,3-Butylene glycol	5.0		
(25) Preservative	0.4	0.4	0.4

*1:

$$CH_{3} \xrightarrow{CH_{3}} \xrightarrow{CH_{3$$

wherein
$$a = 3$$
, $b = 9$, $c = 0$,
 $m = 50$, $n = 10$, $o = 3$
 $R^1 = CH_3$, $R^2 = H$
M.W.: about 6000

Modified Ratio with Polyether: about 20 % by weight

[0033]

Table 2

1

2

3

4

5

į	5		
	_		

10

15

20

25

30

35

40

45

50

55

I	1		9		
(1) Decamethylcyclopentasiloxane	16.5	16.5	55.5	2.5	16.5
(2) Dimethyl polysiloxane	5.0	5.0	5.0	5.0	5.0
(3) Liquid peraffin	3.0	3.0	3.0	3.0	3.0
(4) 2-Ethylhexyl p-methoxycinnamate	1.5	1.5	1.5	1.5	1.5
(5) Dimeticone copolyot*1	2.0	2.0	2.0	2.0	2.0
(6) Sorbitan sesquiisostearate	1.0	1.0	1.0	1.0	1.0
(7) Isostearic acid	0.5	0.5	0.5	0.5	0.5
(8) Solid paraffin	3.0	3.0	3.0	3.0	3.0
(9) Carnauba wax	1.0	1.0	1.0	1.0	1.0
(10) Fragrance	0.1	0.1	0.1	0.1	0.1
(11) Silicone-treated titanium oxide	-	12.0	12.0	12.0	12.0
(12) Titanium oxide	12.0	-	-	-	-
(13) Silicone-treated iron oxide	-	3.0	3.0	3.0	3.0
(14) Iron oxide	3.0	-	•	-	-
(15) Iron oxide-coated titanated mica	0.5	0.5	0.5	0.5	0.5
(16) Silicone-treated sericite	-	3.0	-	3.0	3.0
(17) Sericite	3.0	-	-	-	-
(18) Silicone-treated talc	-	2.5	-	2.5	2.5
(19) Talc	2.5	-	-	-	-
(20) Porous spherical silica (partical size: 4 - 6 μm)	10.0	-	5.0	10.0	-
(21) Non spherical silica (particle size: 4 - 6 μm)	-	-	-	-	10.0
(22) Spherical polymethyl methacrylate	-	10.0	-	-	-
(23) Purified water	30.0	30.0	3.5	45.2	30.0
(24) 1,3-Butylene glycol	5.0	5.0	3.0	5.0	5.0
(25) Preservative	0.4	0.4	0.4	0.4	0.4

*1: See Formula (I) in Table 1, wherein

a = 3, b = 9, c = 0,

Comparative Example

m = 50, n = 10, o = 3, $R^1 = CH_3, R^2 = H$

M.W.: about 6000

Modified Ratio with Polyether: about 20 % by weight

Table 3

	Example			Comparative Examples				
	1	2	3	1	2	3	4	5
Light spreadability	++	++	++	++	++	++	++	±
Freshness	++	++	+	++	+	-	++	+

Table 3 (continued)

	Example			Comparative Examples				
	1	2	3	1	2	3	4	5
Smoothness after application	++	+	+	++	±	+	+	•
Effect in correcting skin roughness	++	++	++	++	++	+	++	-
Long-lasting power	++	++	+	-	++	++	++	++
Fillability in container	++	++	+	+	±	++		+

[0034] As is clear from the results shown in Table 3, Comparative Example 1 did not achieve cosmetic staying power compared with Examples 1 and 2 since it did not contain a hydrophobically treated powder. Further, Comparative Example 2 resulted in the decrease in the smoothness after application and fillability in a container since spherical silica was not contained. Further, Comparative Examples 3 and 4 did not have suitable ratios of the oil and water, and therefore, Comparative Example 3 was inferior in freshness and Comparative Example 4 was inferior in fillability in a container.

[0035] As opposed to the above, Examples 1 and 2 contained suitable amounts of the hydrophobically treated powder, spherical silica, oil, and water, and therefore, all superior in terms of light slip on the skin, freshness at the time of application, smoothness after application, effect of correcting skin roughness, cosmetic staying power after three hours after application, and fillability in a container were obtained. Further, although Example 2 contained spherical PMMA powder in addition to the spherical silica, but the amount blended was suitable, the effect of the present invention was not impaired.

5 Example 4

30

35

40

45

50

55

5

10

[0036] The following components (1) to (17) were used to prepare a cosmetic base as a solid water-in-oil type emulsion cosmetic composition according to the present invention:

	(1) Dimethyl polysiloxane	29.0 wt%
	(2) Methylphenyl polysiloxane	10.0
	(3) Liquid paraffin	5.0
9	(4) Glyceryl trioctanate	5.0
	(5) Microcrystallline wax	2.0
	(6) Carnauba wax	2.0
	(7) Sorbitan sesquiisostearate	3.0
	(8) Polyoxyethylene hydrogenated castor oil	0.5
	(9) Silicone-treated titanium oxide	3.0
	(10) Silicone-treated iron oxide	1.5
	(11) Silicone-treated barium sulfate	4.5
	(12) Boronitride	1.0
	(13) Spherical silica	7.0
	(14) Spherical silicone resin powder	2)
	(15) Purified water	26.0
	(16) Dipropylene glycol	8.0
	(17) Preservative	0.5

Method of Preparation

[0037] The components (1) to (8) were heated to 90°C to melt, then the mixed and pulverized components (9) to (14) were added and stirred to disperse them homogeneously. The homogeneously melted components (15) to (17) were added, then the overall mixture was homogeneously emulsified, then filled in a container and cooled to obtain the solid water-in-oil type emulsion cosmetic base.

Example 5

[0038] The following components (1) to (16) were used to prepare a stick cosmetic composition as a solid water-inoil type emulsion cosmetic composition according to the present invention:

15	(1) Decamethylcyclopentasiloxane	16.0 wt%
	(2) Dodecamethylcyclohexasiloxane	8.0
	(3) Alkyl dimeticone copolyol*1	3.0
20	(4) Solid paraffin	5.0
	(5) Candelilla wax	1.0
	(6) Carnauba wax	1.0
	(7) Silicone-treated titanium oxide	15.0
25	(8) Silicone-treated iron oxide	3.0
	(9) Spherical silica	10.0
	(10) Lauroyllysine powder	2.0
30	(11) Metal soap-treated talc	15.0
	(12) Purified water	10.5
	(13) Sodium hydroxymethoxybenzophenone sulfonate	3.0
	(14) Glycerol	2.0
35	(15) Dipropylene glycol	5.0
	(16) Preservative	0.5

^{*1:} See Formula (I) in Table 1, wherein a=3, b=50, c=60, m=100, n=50, o=15, $R^1=C_{16}H_{33}, R^2=H$

M.W.: about 15000

45 Method of Preparation

40

[0039] The components (1) to (6) were heated to 90°C to melt, then the mixed and pulverized components (7) to (11) were added and stirred to disperse them homogeneously. The homogeneously melted components (12) to (16) were added, then the overall mixture was homogeneously emulsified, then filled in a container and cooled to obtain the solid water-in-oil type emulsion stick cosmetic composition.

Example 6

[0040] The following components (1) to (17) were used to prepare a solid water-in-oil type emulsion foundation according to the present invention:

	(1) Dimethyl polysiloxane	6.0 wt%
5	(2) Decamethylcyclopentasiloxane	22.0
	(3) Dodecamethylcyclohexasiloxane	12.0
	(4) Methylphenyl polysiloxane	5.0
10	(5) Microcrystallline wax	2.0
	(6) Carnauba wax	2.0
	(7) Polyether-modified dimethyl polysiloxane*1	3.0
	(8) Polyoxyethylene hydrogenated castor oil	0.5
15	(9) Silicone-treated titanium oxide	5.0
	(10) Silicone-treated iron oxide	2.5
	(11) Silicone-treated barium sulfate	, 1.5
20	(12) Boronitride	1.0
	(13) Spherical silicone resin powder	2.0
	(14) Spherical silica	7.0
	(15) Purified water	20.0
25	(16) Dipropylene glycol	8.0
	(17) Preservative	0.5
	*1: See Formula (I) in Table 1, wherein	•

M.W.: about 6000

Modified Ratio with Polyether: about 20 % by weight

Method of Preparation

The components (1) to (8) were heated to 90°C to melt, then the mixed and pulverized components (9) to (13) were added and stirred to disperse them homogeneously. The homogeneously dissolved components (15) to (17), in which the component (14) was dispersed, were added, then the overall mixture was homogeneously emulsified, then filled in a container and cooled to obtain the solid water-in-oil type emulsion foundation.

Each of the solid water-in-oil type emulsion cosmetic composition obtained in the above Examples 4 to 6 was evaluated by a panel in the same way as in Example 1. As a result, each was evaluated as being "Excellent" in all items other than the items evaluated as "Good".

[0043] As explained above, the solid water-in-oil type emulsion cosmetic composition according to the present invention is an epochmaking solid water-in-oil type emulsion cosmetic composition not seen in the past which is capable of containing a relatively large amount of water, has a fresh, superior, skin feeling, and effectively corrects skin roughness, while exhibiting a long-lasting power.

Claims 50

30

- 1. A solid water-in-oil type emulsion cosmetic composition comprising a hydrophobically treated powder, spherical silica, an oil, and water, a ratio by weight of the oil to the water (i.e., oil/water) being 0.5 to 10.
- 55 A solid water-in-oil type emulsion cosmetic composition as claimed in claim 1, wherein the amount of the spherical silica blended is 2.0 to 15.0 % by weight and the proportion of the spherical silica in the total spherical powder is 30.0 to 100 % by weight.

a = 3, b = 9, c = 0,

m = 50, n = 10, o = 3,

 $R^1 = CH_3, R^2 = H$

3. A solid water-in-oil type emulsion cosmetic composition as claimed in claim 1, wherein the total amount of the oil

and water is 40.0 to 90.0 % by weight.

5	4.	A solid water-in-oil type emulsion cosmetic composition as claimed in claim 1, wherein the ratio by weight of the oil to the water (i.e., oil/water) is 1 to 5.
	5.	A solid water-in-oil type emulsion cosmetic composition as claimed in claim 1, wherein the cosmetic composition is a make-up cosmetic composition.
10		
15		
20		
25		
30		
35		
40		
45		
50		
55		



Europäisches Patentamt European Patent Office Office européen des brevets



(11) EP 1 036 553 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3: 10.12.2003 Bulletin 2003/50

(51) Int Cl.7: A61K 7/02

(43) Date of publication A2: 20.09.2000 Bulletin 2000/38

(21) Application number: 00105666.2

(22) Date of filing: 17.03.2000

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU

MC NL PT SE

Designated Extension States:

AL LT LV MK RO SI

(30) Priority: 18.03.1999 JP 7443199

(71) Applicant: SHISEIDO COMPANY LIMITED Chuo-ku, Tokyo 104-8010 (JP)

(72) Inventors:

 Daisuke, Aso, c/o Shiseido Research Center Yokohama-shi, Kanagawa 223-8553 (JP)

Akihito, Yokotsuka,
 c/o Shlseido Research Center
 Yokohama-shi, Kanagawa 223-8553 (JP)

(74) Representative: Henkel, Feiler, Hänzel Möhlstrasse 37 81675 München (DE)

(54) Solid water-in-oil type emulsion cosmetic composition

(57) A solid water-in-oil type emulsion cosmetic composition having a fresh, superior skin feeling, and effectively correcting roughness of the skin while exhibiting long-lasting power, which contains a hydrophobi-

cally treated powder, spherical silica, an oil, and water, wherein a ratio by weight of the oil to the water (i.e. oil/water) is 0.5 to 10.



EUROPEAN SEARCH REPORT

Application Number

EP 00 10 5666

Category	Citation of document with inc		Relevant	CLASSIFICATION OF THE
Oziogoi y	of relevant passag	es	to claim	APPLICATION (InLCI.7)
P,X	WO 99 56702 A (FUKU MARI (JP); SHISEIDO 11 November 1999 (19 * claims 1,4,14; tal	CO LTD (JP)) 999-11-11)	1-5	A61K7/02
X	US 5 362 482 A (KUM 8 November 1994 (199 * column 6, line 21 1,4,7; examples 5,6 * column 7, line 37	94-11-08) - line 47; claims ,8.9 *	L) 1,3-5	
A	EP 0 782 881 A (ORE) 9 July 1997 (1997-0) * page 7, line 7 18-22,25,27,37,39;	7-09) line 15; claims	1,3,5	
A	US 5 643 555 A (CANI 1 July 1997 (1997-0 * column 2, line 49 claims 1,3; example	7-01) - column 3, line 37	;	TECHNICAL FIELDS
A	US 5 853 711 A (ITO 29 December 1998 (1 * claim 1; tables I	998-12-29)	1-4	SEARCHED (Int.Cl.7) A61K A61Q
P,A	EP 0 976 388 A (SHI 2 February 2000 (20 * claims 1,2,5,6; e	00-02-02)	1-5	
	The present search report has b	een drawn up for all claims		
	Place of search	Date of completion of the sea	ren	Examiner
	BERLIN	16 October 26	003 An	giolini, D
X:per Y:per doo A tec O:no	ATEGORY OF CITED DOCUMENTS rticularly relevant if taken alone ticularly relevant if combined with anothument of the earne category thrological background newritten disclosure permediate document	T: theory or p E: earlier pat after the fill er D: document L: document	rinciple underlying the ent document, but pub- ing date atted in the application cited for other reasons	r invention Sished on, or n

ANNEX TO THE EUROPEAN SEARCH REPORT ON EUROPEAN PATENT APPLICATION NO.

EP 00 10 5666

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

16-10-2003

Patent document cited in search report	ì	Publication date	l	Patent fami member(s		Publication date
WO 9956702	A	11-11-1999	CN EP WO JP TW US	1272057 1016400 9956702 2000016919 513311 2002022037	A1 A1 A B	01-11-2000 05-07-2000 11-11-1999 18-01-2000 11-12-2002 21-02-2002
US 5362482	Α	08-11-1994	JP JP EP	1079104 2691729 0374332	A B2	24-03-1989 17-12-1997 27-06-1990
EP 0782881	A	09-07-1997	FR DE DE EP ES US	2743004 69600353 69600353 0782881 2120278 5939079	D1 T2 A1 T3	04-07-1997 16-07-1998 10-12-1998 09-07-1997 16-10-1998 17-08-1999
US 5643555	Α	01-07-1997	FR AT CA DE DE EP ES JP JP	2720644 190224 2151012 69515369 69515369 0686391 2145882 2656226 8169808	T A1 D1 T2 A1 T3 B2	08-12-1995 15-03-2000 07-12-1995 13-04-2000 06-07-2000 13-12-1995 16-07-2000 24-09-1997 02-07-1996
US 5853711	Α .	29-12-1998	JP JP JP JP DE DE EP	9095433 3406437 9095434 3406438 9095435 69613647 69613647 0765656 449484	B2 A B2 A D1 T2 A1	08-04-1997 12-05-2003 08-04-1997 12-05-2003 08-04-1997 09-08-2001 08-05-2002 02-04-1997 11-08-2001
EP 0976388	A	02-02-2000	JP EP KR US	2000044425 0976388 2000012102 2001007672	A2 A	15-02-2000 02-02-2000 25-02-2000 12-07-2001